E-Gel[™] 50 bp DNA Ladder

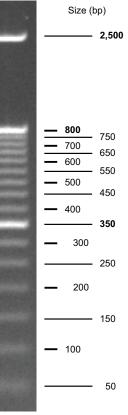
PRODUCT INFORMATION SHEET

Pub. No. MAN0001727

ContentsCatalog No.
10488099Amount
100 applicationsKit contentsStorage• Product is shipped at ambient temperature.
• Store at room temperature or at 4°C for up to 6 months,
or at -20°C for long term storage.• Kit contents

Product description

- The Invitrogen[™] E-Gel[™] 50 bp DNA Ladder is designed for sizing and quantification of double stranded DNA on 2% E-Gel[™] agarose gels.
- The E-Gel[™] 50 bp DNA Ladder consists of 17 individual chromatography-purified DNA fragments ranging in size from 50 bp to 2,500 bp.
- Reference bands at 350 bp, 800 bp, and 2,500 bp are included for easy orientation.
- The ladder is supplied with 1X E-Gel[™] Sample Loading Buffer for sample DNA.



Rev. A.0

Visit our product pages for additional information and protocols.
 Online
 Co opling to view related DNA ladders and markers

resources

- Go online to view related DNA ladders and markers.
- For support, visit thermofisher.com/support.

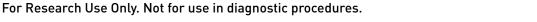


Required materials

- E-Gel[™] E-Gel[™] EX or E-Gel[™] Agarose Gel with SYBR[™] Safe (See **Choosing the** right DNA ladder for your E-Gel[™] agarose gel)
- TE Buffer (Cat. No. AM9858)
- Ultrapure[™] DNase/RNase-Free Distillated Water (Cat. No. 10977023)

Important guidelines

- Do not heat the E-Gel[™] 50 bp DNA Ladder before loading.
- Load the same volume of DNA sample and DNA ladder.
- For quantification, adjust the concentration of the sample to equalize it approximately with the amount of DNA in the nearest band of the ladder.
- Dilute sample DNA in TE buffer to avoid degradation of DNA sample.
- Choosing the right DNA ladder for your E-Gel™ agarose gel
- Troubleshooting
- Limited product warranty and disclaimer details





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Prepare DNA ladders and samples for electrophoresis

This protocol provides a brief description of how to use the DNA ladder with E-Gel[™] agarose gels. For detailed instructions on using specific types of E-Gel[™] agarose gels, go to thermofisher.com or contact Technical Support.

	Step		Action
1		Prepare DNA ladder	 a. Thaw, mix and briefly centrifuge DNA ladder before use. b. Prepare DNA ladder. For 2% E-Gel[™] EX Agarose Gels, mix 2 µL of DNA ladder with 18 µL of water. For 2% E-Gel[™] Agarose Gels, mix and use the ladder without dilution. For 2% E-Gel[™] 48 Agarose Gels, mix 2 µL of DNA ladder with 13 µL of water.
2		Prepare samples	a. Dilute your sample 2- to 10-fold with TE Buffer (Cat. No. AM9858), 1X E-Gel™ Sample Loading Buffer (Cat No. 10482055), or water. b. Mix gently.
3		Load samples and DNA ladders	 a. Load DNA ladders and DNA samples into the appropriate wells of the E-Gel[™] agarose gel. Add 20 μL for E-Gel[™] and E-Gel[™] EX Agarose Gels. Add 15 μL for E-Gel[™] 48 Agarose Gels. b. Add water to any empty wells, so that all wells contain an equal volume of liquid.
		Perform electrophoresis	a. Choose the appropriate E-Gel [™] run protocol for your gel type based on the electrophoresis device being used.
			Gel type Program Recommended run time
			E-Gel [™] Power Snap Electrophoresis Device (Cat. No. G8100)
4			E-Gel [™] EX Agarose Gel (2%) E-Gel EX 4 1-2% 15 min (20 min max)
			E-Gel [™] Agarose Gel (2%) E-Gel 0.8-2% 26 min (40 min max)
			E-Gel [™] E-Base [™] Device
			E-Gel [™] 48 Agarose Gel (2%) EG 20 min
			b. Run the program to start electrophoresis.
5		Visualize agarose gel	Visualize DNA ladder and samples.
			 Use the E-Gel[™] Power Snap Camera (Cat. No. G8200), E-Gel[™] Imager (Cat. No. 466612), or other blue light imager to detect DNA bands stained with SYBR[™] stains.
			 UV transilluminator to detect DNA bands stained with ethidium bromide.

For support, visit thermofisher.com/support.

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